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Patentanmeldung Nr. Patent application No. Demande de brevet n°

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Der Präsident des Europäischen Patentamts;
Im Auftrag

For the President of the European Patent Office

Le Président de l'Office européen des brevets
p.o.

R C van Dijk



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(Falls die Bezeichnung der Erfindung nicht angegeben ist, siehe Beschreibung.
If no title is shown please refer to the description.
Si aucun titre n'est indiqué se référer à la description.)

Process for the preparation of enantiomerically pure 1-substituted-3-aminoalcohols

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Description

The present invention refers to a process for the preparation of enantiomerically pure 1-substituted-3-aminoalcohols, particularly of (*S*)-(-)- and (*R*)-(+)-3-*N*-methylamino-1-(2-thienyl)-1-propanol, which can be obtained by asymmetrically hydrogenating salts of the corresponding aminoketones and a carboxylic acid, particularly of 3-*N*-methylamino-1-(2-thienyl)-1-propanone and a carboxylic acid, in the presence of a catalyst comprising a transition metal and a diphosphine ligand.

(*S*)-(-)-3-*N*-Methylamino-1-(2-thienyl)-1-propanol is an intermediate for the preparation of (*S*)-(+)-methyl-[3-(1-naphthyloxy)-3-(2-thienyl)-propyl]-amine (duloxetine), an agent for the treatment of depression and urinary incontinence (Huiling et al. *Chirality* 2000, 12, 26-29, Sorbera et al. *Drugs of the Future* 2000, 25(9), 907-916).

Several processes for racemic (WO4/005239) and asymmetric (Sorbera et al. below) hydrogenation of thienyl amino ketone are known, as well as processes for chiral resolution of the resulting 3-*N*-methylamino-1-(2-thienyl)-1-propanol (WO-A 2004/005220, WO-A 2004/005307). Furthermore, processes for direct asymmetric hydrogenation using transition metal-ligand complexes are disclosed in EP-A 0 647 648, EP-A 0 926 152, EP-A 0 945 457, EP-A 0 955 303 and WO-A 02/40492.

Huiling et al. describe a preparation of (*S*)-(-)-3-*N*-methylamino-1-(2-thienyl)-1-propanol from thiophene. Thiophene is converted with 3-chloropropanoyl chloride in the presence of tin tetrachloride in benzene to 3-chloro-1-(2-thienyl)-1-propanone, which is reduced with sodium borohydride in ethanol to 3-chloro-1-(2-thienyl)-1-propanol. Kinetic resolution by transesterification using vinyl butanoate and lipase B from *Candida antarctica* as catalyst in hexane yielded (*S*)-3-chloro-1-(2-thienyl)-1-propanol, which is converted to (*S*)-3-iodo-1-(2-thienyl)-1-propanol using sodium iodide in acetone. Subsequent treatment with methylamine in tetrahydrofuran afforded (*S*)-(-)-3-*N*-methylamino-1-(2-thienyl)-1-propanol.

Sorbera et al. describe another preparation of (*S*)-(-)-3-*N*-methylamino-1-(2-thienyl)-1-propanol from thiophene, which is essentially the same as the one described by Huiling et al. except that 3-chloro-1-(2-thienyl)-1-propanone is asymmetrically reduced to (*S*)-3-chloro-

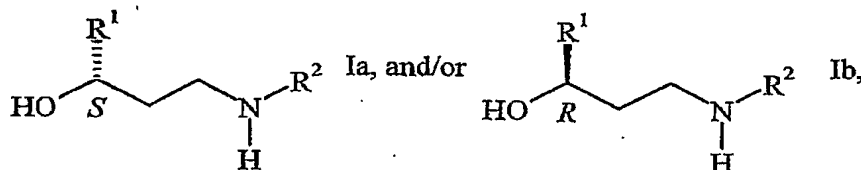
1-(2-thienyl)-1-propanol using borane and catalytic amounts of (*R*)-3,3-diphenyl-1-methyltetrahydro-3*H*-pyrrolo[1,2-*c*][1,3,2]oxazaborole in THF. This asymmetric reduction afforded (*S*)-3-chloro-1-(2-thienyl)-1-propanol in a yield of 86% from 3-chloro-1-(2-thienyl)-1-propanone (Wheeler et al. *J. Label. Compd. Radiopharm.* 1995, 36, 213-223).

The drawbacks of the preparations of (*S*)-(-)-3-*N*-methylamino-1-(2-thienyl)-1-propanol above, are the use of toxic or carcinogenic compounds such as tin tetrachloride and benzene and/or the use of expensive compounds such as borane or sodium iodide, the latter being in addition difficult to dispose of. The disclosed asymmetric hydrogenation processes with diphosphines are not satisfying in regard of the hydrogenation of 3-*N*-methylamino-1-(2-thienyl)-1-propanone.

It is an object of the present invention to provide an ecological and economical process for the preparation of enantiomerically pure 1-substituted-3-aminoalcohols, particularly of (*S*)-(-)- and (*R*)-(+)-3-*N*-methylamino-1-(2-thienyl)-1-propanol. It is another object of the present invention to provide new salts of 3-*N*-methylamino-1-(2-thienyl)-1-propanone and organic acids.

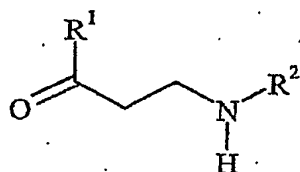
These objects are achieved by the process of claim 1.

Provided is a process for the preparation of salts of a carboxylic acid with an aminoalcohol of the formula



wherein R^1 is selected from the group consisting of 2-thienyl, 2-furanyl and phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups, and wherein R^2 is C_{1-4} -alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups, comprising asymmetrically hydrogenating a salt of a carboxylic acid with an aminoketone of the formula

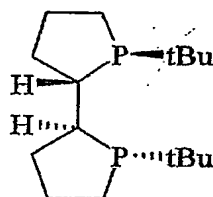
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II,

5 wherein R^1 and R^2 are as defined above,
in the presence of a transition metal complex of a diphosphine ligand.

Furthermore provided is a process for the preparation of salts of a carboxylic acid with an aminoalcohol of the formula Ia or Ib, wherein R^1 is selected from the group consisting of
10 2-thienyl, 2-furanyl and phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups, and wherein R^2 is C_{1-4} -alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups,
comprising asymmetrically hydrogenating a salt of a carboxylic acid with an aminoketone of
15 the formula II, wherein R^1 and R^2 are as defined above, in the presence of a transition metal complex of an aryl- or biaryl-diphosphine ligand or the diphosphine ligand



(R,R,S,S)-TangPhos.

Asymmetric hydrogenation of 1-substituted-3-N-alkylamino-1-propanone hydrochlorides with
25 diphosphine-transition metal complexes is not possible without generating the free amines by neutralizing the acidic salts. Due to the fact that the free 1-substituted-3-N-alkylamino-1-propanones tend to decompose, the resulting 1-substituted-3-N-alkylamino-1-propanols are contaminated with by-products. Exchanging the chloride anion by a carboxylic acid allows direct hydrogenation of the resulting, optionally purified, salt of such acid with a 1-substituted-
30 3-N-alkylamino-1-propanone in high yields, high purity and high enantiomeric excess (ee). Avoiding decomposition of the free aminoketone in the presence of base is another advantageous feature of the present invention.

Carboxylic acids in the meaning of the present invention are carboxylic acids having one free carboxyl group which can form a salt with an amino compound of formulae II and/or I.

Particularly preferred the carboxylic acids are monocarboxylic acids. Dicarboxylic or tri-carboxylic acids which do not form an inner salt like fumaric, maleic or adipic acid tend to give unusable resinous precipitates. However, carboxylic acids which form an inner salt and still have one free carboxy group should be comprised in the definition of carboxylic acids in the meaning of the present invention. Examples of said carboxylic acids having more than one carboxy group, but having only one free carboxy group, are amino acids, such as aspartic acid or glutamic acid.

In a preferred process, the carboxylic acid is selected from the group consisting of optionally substituted C₁₋₁₈-alkanoic acids and optionally substituted mono- and bicyclic aromatic acids.

In a preferred embodiment the carboxylic acids are optionally substituted with one or more C₁₋₆-alkyl, C₁₋₆-alkoxy, aryl, amino, optionally protected carbonyl, halogen or hydroxy groups and optionally further carboxylic groups.

Examples for C₁₋₁₈-alkanoic acids in the meaning of the present process are butyric acid, valeric acid, caproic acid, pelargonic acid, lauric acid, palmitic acid, stearic acid, 2-hydroxybutyric acid, 3-hydroxy-3-methylbutyric acid, 2-hydroxy-4-phenylbutyric acid, L-aspartic acid, D-aspartic acid, DL-aspartic acid, 2-keto-L-gulonic acid, 2-keto-D-gulonic acid, (-)-2,3:4,6-di-O-isopropylidene-2-keto-L-gulonic acid and (+)-2,3:4,6-di-O-isopropylidene-2-keto-D-gulonic acid.

In a further preferred process the carboxylic acid is a mono- and bicyclic aromatic acid, optionally substituted with one or more C₁₋₆-alkyl, C₁₋₆-alkoxy, halogen or hydroxy groups.

Examples for mono- and bicyclic aromatic carboxylic acids in the meaning of the present process are benzoic acid, salicylic acid, 3-methyl-benzoic acid and 1-, 2- or 3-naphthalene carboxylic acid.

Here and hereinbelow the term "enantiomerically pure compound" comprises optically active compounds with an enantiomeric excess (ee) of at least 90 %.

Here and hereinbelow the term "C_{1-n}-alkyl" represents a linear or branched alkyl group having 1 to n carbon atoms. Optionally with one or more halogen atoms substituted C₁₋₆-alkyl represents for example methyl, ethyl, propyl, isopropyl, butyl, isobutyl, *sec*-butyl, *tert*-butyl, pentyl and hexyl.

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Here and hereinbelow the term "C_{1-n}-alkoxy" represents a linear or branched alkoxy group having 1 to n carbon atoms. Optionally with one or more halogen atoms substituted C₁₋₆-alkoxy represents for example methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, *sec*-butoxy, *tert*-butoxy, pentyloxy and hexyloxy.

10

Here and hereinbelow the term "C_{3-n}-cycloalkyl" represents a cycloaliphatic group having 3 to n carbon atoms. Optionally with one or more halogen atoms substituted C₃₋₁₀-cycloalkyl represents for example mono- and polycyclic ring systems such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, adamantyl or norbornyl.

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Here and hereinbelow the term "C_{3-n}-cycloalkoxy" represents a cycloalkoxy group having 3 to n carbon atoms. Optionally with one or more halogen atoms substituted C₃₋₁₀-cycloalkyl represents for example cyclopropoxy, cyclobutoxy, cyclopentyloxy, cyclohexyloxy, cycloheptyloxy, cyclooctyloxy and cyclodecyloxy.

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Here and hereinbelow the term "aryl" represents an aromatic group, preferably phenyl or naphthyl, optionally being substituted with one or more halogen atoms, nitro and/or amino groups, and/or optionally substituted C₁₋₆-alkyl, C₁₋₆-alkoxy or di-C₁₋₆-alkylamino groups, wherein the alkyl moieties optionally are substituted with one or more halogen atoms.

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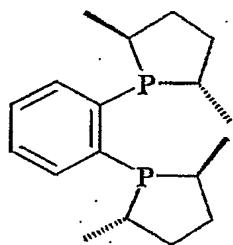
In a preferred embodiment R¹ is 2-thienyl, optionally substituted with one or more halogen atoms, and R² is methyl or ethyl.

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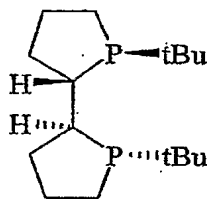
In a further preferred embodiment, the compound of formula II is selected from the group consisting of (*S*)-(-)-3-*N*-methylamino-1-(2-thienyl)-1-propanol, (*S*)-(-)-3-*N*-methylamino-1-(3-chloro-2-thienyl)-1-propanol, (*R*)-(+)-3-*N*-methylamino-1-(2-thienyl)-1-propanol and (*R*)-(+)-3-*N*-methylamino-1-(3-chloro-2-thienyl)-1-propanol.

In a preferred process, the transition metal is selected from the group consisting of rhodium, ruthenium and iridium, preferably rhodium.

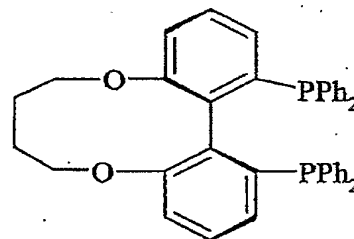
In a further preferred process, the diphosphine ligand is selected from the group consisting of



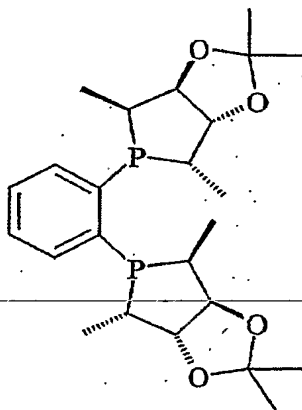
(S,S)-Me-DuPhos,



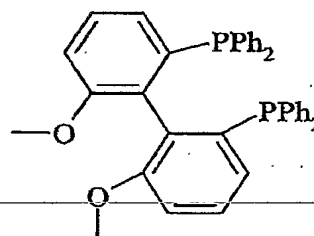
(R,R,S,S)-TangPhos



(S)-C4-TunaPhos,



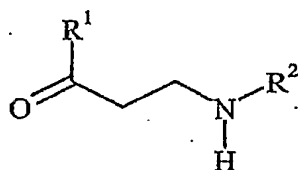
(S,S,S,S)-Me-KetalPhos



(S) and (R)-MeO-BiPhep.

In a preferred process, the compounds of formulae Ia and Ib are obtained from their corresponding salts with a carboxylic acid by aqueous hydrolysis in the presence of an alkali or earth alkali hydroxide.

Provided are salts of a carboxylic acid with an aminoketone of the formula



II,

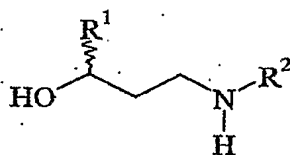
wherein R^1 is selected from the group consisting of 2-thienyl, 2-furanyl and phenyl, each

optionally substituted with one or more halogen atoms and/or one or more C₁₋₄-alkyl or C₁₋₄-alkoxy groups, and wherein R² is C₁₋₄-alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more C₁₋₄-alkyl or C₁₋₄-alkoxy groups.

- 5 Particularly preferred, the salt contains a carboxylic acid selected from the group consisting of C₁₋₁₈-alkanoic acids, (-)-2,3:4,6-di-*O*-isopropylidene-2-keto-L-gulonic acid, (+)-2,3:4,6-di-*O*-isopropylidene-2-keto-D-gulonic acid, 2-keto-L-gulonic acid, 2-keto-D-gulonic acid, L-aspartic acid, D-aspartic acid, benzoic acid, 3-methyl-benzoic acid, salicylic acid and 2-naphthalene carboxylic acid

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Furthermore provided are salts of a carboxylic acid with an aminoalcohol of the formula



I,

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- wherein R¹ is selected from the group comprising 2-thienyl, 2-furanyl and phenyl, each optionally substituted with one or more halogen atoms and/or one or more C₁₋₄-alkyl or C₁₋₄-alkoxy groups, and wherein R² is C₁₋₄-alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more C₁₋₄-alkyl or C₁₋₄-alkoxy groups, with the exception of salts, wherein the acid is (-)-2,3:4,6-di-*O*-isopropylidene-2-keto-L-gulonic acid or (+)-2,3:4,6-di-*O*-isopropylidene-2-keto-D-gulonic acid.

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The present invention is illustrated by the following non-limiting examples.

25 Examples

Example 1: Preparation of 3-*N*-methylamino-1-(2-thienyl)-1-propanone hydrochloride (PRON₁MT2-HCl)

- A mixture of 2-acetylthiophene (25.5 g, 200 mmol), methylamine hydrochloride (14.9 g, 220 mmol), paraformaldehyde (8.2 g, 280 mmol) and ethanol (100 mL) is heated in an autoclave at 120 to 130 °C for 9 hours. The obtained light brown solution is cooled to 20 °C and part of the ethanol (50 mL) is removed by distillation in vacuo. Ethyl acetate (200 mL) is added to the residue to afford a thick suspension, which is cooled to 0 °C and kept for 45 min at

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that temperature. The obtained precipitate is isolated by filtration and dried, yielding 29.3 g 3-*N*-methylamino-1-(2-thienyl)-1-propanone hydrochloride (PRON,MT2-HCl, 71%) as a slightly yellow powder.

5 **Comparative example 1: Preparation of racemic 3-*N*-methylamino-1-(2-thienyl)-1-propanol (PROL,MT)**

Sodium hydroxide (4.0 g of a 50% aqueous solution) is added to a mixture of PRON,MT2-HCl (10.3 g, 50 mmol) and ethanol (35 mL) at 4 °C in about 5 min. Neat sodium borohydride (0.95 g, 25 mmol) is added in several portions in about 30 min to afford a beige suspension which is stirred at 4 °C for additional 4 hours. Acetone (10 mL) is added dropwise in 5 min and the mixture is stirred for additional 10 min before water (20 mL) is added. The mixture is concentrated about 5 times in vacuo and the obtained residue is extracted with *tert*-butyl methyl ether (MTBE) (2 × 20 mL). The collected organic phases are concentrated in vacuo affording 7.2 g racemic 3-*N*-methylamino-1-(2-thienyl)-1-propanol (PROL,MT, 84%) as an orange oil which crystallizes spontaneously after a few hours. ¹H-NMR (DMSO-*d*₆, 400 MHz): 7.35 (1 H, dd, *J* = 4.8, 1.0), 6.94 (1 H, dd, *J* = 4.8, 3.6), 6.90 (1 H, dd, *J* = 3.6, 1.0), 4.90 (1 H, t), 3.7 (2 H, m), 2.56 (2 H, m), 2.25 (3 H, s), 1.79 (2 H, q); ¹³C-NMR (DMSO-*d*₆, 100 MHz): 150.9, 126.3, 123.7, 122.3, 67.8, 48.5, 38.7, 36.0.

20 **Comparative example 2: Preparation of (*S*)-(-)-3-*N*-methylamino-1-(2-thienyl)-1-propanol (PROL,MT,S)**

In a 50 mL autoclave a solution of PRON,MT2-HCl (250 mg, 0.56 mmol) in methanol (5 mL) and an equivalent amount of NaOH mixture is charged under nitrogen. Afterwards, a solution of [Rh((*S,S*)-Me-Duphos)]BF₄ (2.7 mg) in methanol (2 mL) prepared under nitrogen is added via a syringe. The autoclave is then closed and purged several times with nitrogen, heated up to 50 °C, then hydrogen is added until the pressure reaches 30 bar. After 5 hours at that temperature under stirring, the autoclave is cooled to room temperature. Once cold, the clear yellow-brownish solution is transferred into a 50 mL round bottom flask and concentrated to dryness affording a beige solid (0.23 g, 92%, ee: about 97% by HPLC).

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Comparative example 3: Preparation of PROL,MT,S

A solution of PRON,MT2-HCl (250 mg, 0.56 mmol) in methanol (5 mL) is charged under nitrogen in a 50 mL autoclave. Afterwards, a solution of 1.8 mg Rh(cod)₂BF₄ and 2.7 mg the

ligand of formula III, with $R^3 = \text{OMe}$, $R^4 = R^5 = \text{dicyclohexylphosphinyl}$ and $R^6 = R^7 = \text{diphenylphosphinyl}$, in methanol (2 mL) previously prepared by stirring the components for 15 min under nitrogen is added via a syringe. The hydrogenation is carried out as described above, affording PROL,MT,S as a beige solid (0.21 g, 84%, ee: about 11% by HPLC)

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Example 2: Preparation of 3-N-methylamino-1-(2-thienyl)-1-propanone and its (-)-2,3:4,6-di-O-isopropylidene-2-keto-L-gulonic acid salt (PRON,MT2-diketegulac)

A mixture of PRON,MT2-HCl (15.0 g, 47.5 mmol), MTBE (170 mL) and water (20 mL) is cooled to 0 °C, then sodium hydroxide (12.8 g of a 20% aqueous solution) is added dropwise in 15 min and stirred for 10 min. Afterwards, the stirring is stopped, the two phases are separated and the organic one is washed with water (60 mL). Then the collected aqueous phases are extracted with MTBE (2 × 50 mL). To the two collected organic layers is then added dropwise a solution of (-)-2,3:4,6-di-O-isopropylidene-2-keto-L-gulonic acid ((-)-DAG, 13.1 g, 4.48 mol) in MTBE (400 mL). The product precipitates already during addition. At the end of the addition, the suspension is concentrated in vacuo to half the volume, the residue is heated to 50 to 60 °C and heptane (250 mL) is added. Afterwards, the suspension is cooled down to 5 °C, the precipitate is filtered off and washed with MTBE/heptane (1:2, v:v, 2 × 60 mL). Drying at 30 °C for 15 hours at 30 mbar affords a white solid (13.2 g, 66%). (assay is 96.1 weight%, purity is 99.6 area%, by HPLC)

20

Example 3: Preparation of PRON,MT2-diketegulac

A sticky suspension of 675 g (2.13 mol) PRON,MT2-HCl in 5.5 L MTBE and 0.9 L water is stirred at a temperature of 0 to 5 °C in a 10 L vessel. Within 0.5 hours 576 g (2.88 mol) of a 20% NaOH solution are added and the reaction mixture is stirred for another 0.5 hours at the same temperature. After phase separation and washing of the organic phase with water (1.3 L) and extracting the aqueous phase with MTBE (2 × 1.3 L), the combined organic layers are cooled in a 10 L vessel to below 10 °C. After addition of a solution of 590 g (-)-DAG (2.02 mol) within 15 min a yellowish mixture is obtained and crystallization occurs spontaneously or after addition of a crystallization aid, such as a small crystal of the product. After further stirring for 2 hours at 0 to 5 °C, filtration, washing with MTBE (2 × 1.5 L) and drying in in vacuo at 50 to 55 °C, the product (740 g of white-tan solid matter) is obtained.

Example 4: Recrystallization of PRON,MT2-diketegulac

A suspension of 738 g of the product of example 3, 5.3 L MTBE and 2.7 L methanol are heated under reflux. After addition of further 1.8 L methanol at the same temperature a clear yellowish solution is obtained. During cooling to 0 to 5 °C within 3 hours, the product precipitates. After further stirring at 0 to 5 °C for 2 hours, filtration, washing with MTBE (2 × 1 L) and drying in in vacuo at 50 to 55 °C, the product (538 g of white solid) is obtained.

Example 5: Hydrogenation of PRON,MT2-diketegulac

A solution of PRON,MT2-diketegulac of example 4 (250 mg, 0.56 mmol) in methanol (5 mL) is charged under nitrogen in a 50 mL autoclave. Afterwards, a solution of 3 mg [Rh((R,R,S,S)-Tangphos)(norbornadiene)]BF₄ in methanol (2 mL) is added via a syringe to the first mixture. The hydrogenation is carried out as described above, affording the salt of (S)-(-)-3-N-methylamino-1-(2-thienyl)-1-propanol and (-)-DAG (PROL,MT,S-diketegulac) as a beige solid (0.22 g, 92%). Conversion is 100% by HPLC, ee is 95% (S isomer id preferably formed).

Example 6: Hydrolysis of (S)-PROL,MT-diketegulac

9.0 g (20.2 mmol) of solid PROL,MT-diketegulac of example 5 is added in several portions to a mixture of water (22 mL), CH₂Cl₂ (18 mL) and an aqueous solution of sodium hydroxide (30%, 2.07 g, 25.9 mmol) and the reaction mixture (two phases) is stirred for 15 min. After phase separation, the aqueous phase is extracted with CH₂Cl₂ (12 mL). The combined organic phases are washed with water (13 mL). The solvent is partly removed at 20 °C in vacuo to a volume of 9 mL. Heptane (18 mL) is added to the residue and the resulting solution is further concentrated to about 18 mL in vacuo at 20 °C. Crystallization occurs spontaneously or after seeding and the suspension is further stirred for 30 min at 20 °C. The precipitate is filtrated, washed with heptane (7 mL) and dried at 40 °C for 15 hours at 25 mbar affording PROL,MT,S (3.0 g of white solid, 87%).

Example 7: Preparation of PRON,MT2-2-keto-L-gulonate

A mixture of 15.0 g (47.5 mmol) PRON,MT2-HCl, 170 mL MTBE and 20 mL water is cooled to 5 to 10 °C in a 250 mL vessel. After addition of 12.8 g of a 20% NaOH solution the mixture is stirred for further 15 min, while phase separation occurs. The organic phase is washed with water (60 mL) and the aqueous phase is extracted with MTBE (2 × 50 mL). The combined organic phases are cooled to below 10 °C and within 15 min a suspension of 8.7 g (45 mmol)

2-keto-L-gulonic acid in 400 mL MTBE is added. A sticky suspension is formed. The volume of the solvent is reduced to approximately the half. The reaction mixture is heated under reflux and 250 mL heptane are added. After further addition of 300 mL methanol and heating under reflux for 30 min, the mixture is cooled to room temperature (RT) and the solvent is removed.
5 The resinous residue is mixed with 100 mL methanol and the solid matter is filtered off and dried in in vacuo at 50 to 55 °C to yield 8.3 g tan solid matter.

Example 8: Recrystallization of PRON,MT2-2-keto-L-gulonate

8.2 g solid product of example 7 is heated under reflux with 100 mL ethanol. Under stirring, the
10 clear solution is cooled to RT and a resin deposits. The mixture is stirred for 1 hour at RT. The moist resin is dried in in vacuo at 50 to 55 °C to afford 4.5 g of tan solid matter.

Example 9: Preparation of PRON,MT2-benzoate

A mixture of 15.0 g (47.5 mmol) PRON,MT2-HCl, 170 mL MTBE and 20 mL water is cooled
15 to 5 to 10 °C in a 250 mL vessel. After addition of 12.8 g of a 20% NaOH solution the mixture is stirred for further 15 min, while phase separation occurs. The organic phase is washed with water (60 mL) and the aqueous phase is extracted with MTBE (2 × 50 mL). The combined organic phases are cooled to below 10 °C and within 15 min a suspension of 5.5 g (45 mmol) benzoic acid in 400 mL MTBE is added. An oily suspension is formed. The volume of the
20 solvent is reduced to approximately the half. The reaction mixture is heated under reflux and 250 mL heptane are added. The mixture is cooled to RT and stirred for 1 hour. The solid matter is filtered off, washed with Heptane/MTBE (2 × 60 mL) and dried in in vacuo at 50 to 55 °C. (Yield: 10.4 g yellow-brown solid product)

Example 10: Recrystallization of PRON,MT2-benzoate

10.3 g solid product of example 9 is heated under reflux with 50 mL ethylacetate. Under stirring, the clear solution is cooled to RT and a solid deposits. The mixture is stirred for 1 hour at RT. The resin is dried in in vacuo at 50 to 55 °C which affords 6.1 g of light tan solid matter.

Example 11: Hydrogenation of PRON,MT2-benzoate

30 A solution of PRON,MT2-benzoate of example 10 (146 mg) in methanol (5 mL) is charged under nitrogen in a 50 mL autoclave. Afterwards, a solution of 3 mg [Rh((S,S)-Me-Duphos)(1,4-cyclooctadiene)]BF₄ in methanol (2 mL) is added via a syringe. The mixture is

hydrogenated as described above affording 0.12 g solid product (PRON,MT,S-benzoate).
Conversion is 99% by HPLC, ee is 96.7% (*S* isomer preferably formed).

Comparative example 4: Preparation of PRON,MT2-*p*-toluenesulfonate

- 5 A mixture of 15.0 g (47.5 mmol) PRON,MT2-HCl, 170 mL MTBE and 20 mL water is cooled to 5 to 10 °C in a 250 mL vessel. After addition of 12.8 g of a 20% NaOH solution the mixture is stirred for further 15 min, while phase separation occurs. The organic phase is washed with water (60 mL) and the aqueous phase is extracted with MTBE (2 × 50 mL). The combined organic phases are cooled to below 10 °C and within 15 min a suspension of 8.6 g (45 mmol)
- 10 *p*-toluenesulfonic acid monohydrate in 400 mL MTBE is added. An oily suspension is formed. The volume of the solvent is reduced to approximately the half. The reaction mixture is heated under reflux and 250 mL heptane are added. The mixture is cooled to RT and stirred for 30 min. The solid matter is filtered off, washed with MTBE (2 × 60 mL) and dried in in vacuo at 50 to 55 °C affording 14.3 g of product (tan solid matter).

15

Comparative example 5: Recrystallization of PRON,MT2-*p*-toluenesulfonate

- 14.2 g solid product of comparative example 4 is heated under reflux with 50 mL isopropanol. Under stirring, the clear solution is cooled to RT and a solid deposits. The mixture is stirred for 1 hour at RT. The resin is dried in in vacuo at 50 to 55 °C which affords 12.5 g of tan solid
- 20 matter.

Comparative example 6: Hydrogenation of PRON,MT2-*p*-toluenesulfonate

- A solution of PRON,MT2-*p*-toluenesulfonate of comparative example 5 (155 mg) in methanol (5 mL) is charged under nitrogen in a 50 mL autoclave. Afterwards, a solution of 3 mg
- 25 [Rh(Me-Duphos)(1,4-cycloocatadiene)]BF₄ in methanol (2 mL) is added via a syringe. The hydrogenation is carried out as described above, affording 0.12 g solid product (PRON,MT,S-*p*-toluenesulfonate). Conversion is 5% by HPLC, ee is >90% (*S* isomer preferably formed).

Example 12: Preparation of PRON,MT2-laurate

- 30 A mixture of 15.0 g (47.5 mmol) PRON,MT2-HCl, 170 mL MTBE and 20 mL water is cooled to 5 to 10 °C in a 250 mL vessel. After addition of 12.8 g of a 20% NaOH solution the mixture is stirred for further 15 min, while phase separation occurs. The organic phase is washed with water (60 mL) and the aqueous phase is extracted with MTBE (2 × 50 mL). The combined

organic phases are cooled to below 10 °C and within 15 min a suspension of 9.0 g (45 mmol) dodecanoic acid in 200 mL MTBE is added. After 1 hour stirring no product has solidified. The solvent is removed in in vacuo and the oily residue is solved in 50 mL acetonitrile and heated under reflux. The mixture is cooled to RT and stirred for 30 min. At first an oil secretes, which crystallizes at a temperature of below 30 °C. The suspension is further stirred for 30 min at RT, than 30 mL acetonitrile are added to the thickened suspension. The solid matter is filtered off, washed with cold acetonitrile (2 × 10 mL) and dried in in vacuo at below 30 °C, affording 10.9 g product (10.9 g white solid matter).

Example 13: Recrystallization of PRON,MT2-laurate

10.7 g solid product of example 12 is heated under reflux with 70 mL acetonitrile. Under stirring, the clear solution is cooled to RT and an oil secrets, which crystallizes at below 30 °C. The mixture is stirred for 30 min at 10 to 15 °C and 30 mL acetonitrile are added to the thickened suspension. The solid matter is filtered off, washed with cold acetonitrile (2 × 20 mL) and dried in in vacuo at below 30 °C. Yield: 6.3 g white solid matter.

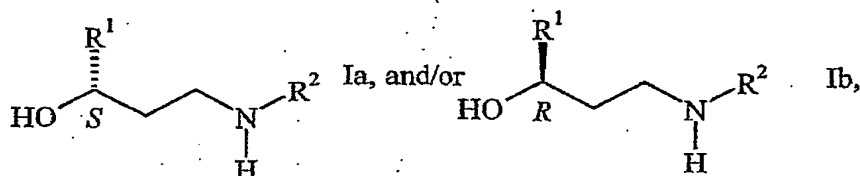
Example 14: Hydrogenation of PRON,MT2-laurate

A solution of PRON,MT2-laurate of example 13 (184 mg) in methanol (5 mL) is charged under nitrogen in a 50 mL autoclave. Afterwards, a solution of 3 mg [Rh((R,R,S,S)-Tang-phos)(norbornadiene)]BF₄ in methanol (2 mL) is added via a syringe. The mixture is hydrogenated as described above, affording 0.16 g solid product (PROL,MT,S-laurate). Conversion is 100% by HPLC, ee is 93.6% (S isomer preferably formed).

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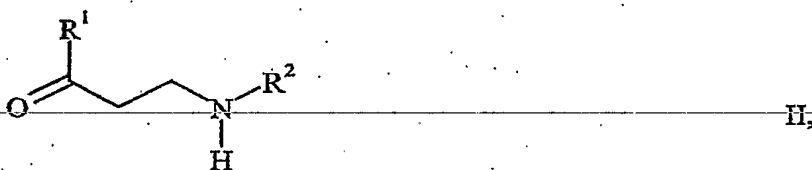
Claims

1. A process for the preparation of salts of
a carboxylic acid with an aminoalcohol of the formula



10 wherein R^1 is selected from the group consisting of 2-thienyl, 2-furanyl and phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups, and wherein R^2 is C_{1-4} -alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups,

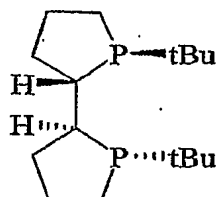
- 15 comprising asymmetrically hydrogenating a salt of
a carboxylic acid with an aminoketone of the formula



wherein R^1 and R^2 are as defined above,
in the presence of a transition metal complex of a diphosphine ligand.

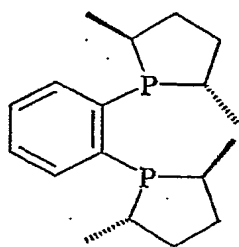
2. A process for the preparation of salts of a carboxylic acid with an aminoalcohol of the
25 formula Ia or Ib, wherein R^1 is selected from the group consisting of 2-thienyl, 2-furanyl and phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups, and wherein R^2 is C_{1-4} -alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups,
30 comprising asymmetrically hydrogenating a salt of a carboxylic acid with an aminoketone of the formula II, wherein R^1 and R^2 are as defined above, in the presence of a transition

metal complex of an aryl- or biaryldiphosphine ligand or the diphosphine ligand

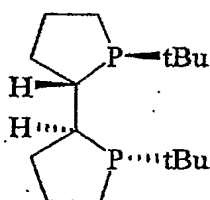


(*R,R,S,S*)-TangPhos.

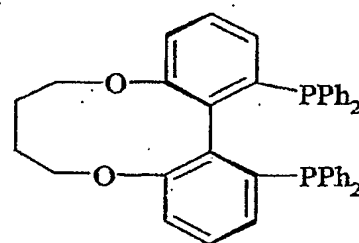
3. The process of claim 1 or 2, wherein the carboxylic acid is selected from the group consisting of optionally substituted C₁₋₁₈-alkanoic acids and optionally substituted mono- and bicyclic aromatic acids.
4. The process of any of claim 1 to 3, wherein R¹ is 2-thienyl, optionally substituted with one or more halogen atoms, and R² is methyl or ethyl.
5. The process of claim 4, wherein the compound of formula II is selected from the group consisting of (*S*)-(-)-3-*N*-methylamino-1-(2-thienyl)-1-propanol, (*S*)-(-)-3-*N*-methylamino-1-(3-chloro-2-thienyl)-1-propanol, (*R*)-(+)-3-*N*-methylamino-1-(2-thienyl)-1-propanol and (*R*)-(+)-3-*N*-methylamino-1-(3-chloro-2-thienyl)-1-propanol.
6. The process of any of claims 1 to 5, wherein the transition metal is selected from the group consisting of rhodium, ruthenium or iridium, and preferably is rhodium.
7. The process of any of claims 1 to 6, wherein the diphosphine ligand is selected from the group consisting of



(*S,S*)-Me-DuPhos,

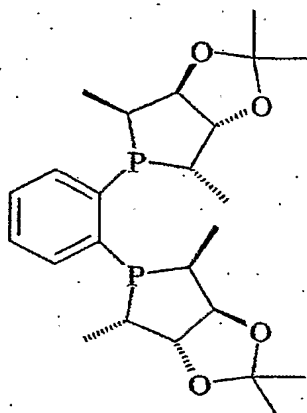


(*R,R,S,S*)-TangPhos

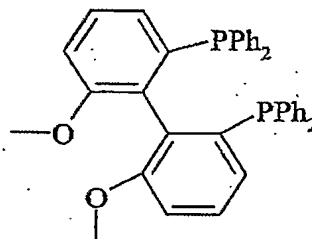


(*S*)-C4-TunaPhos,

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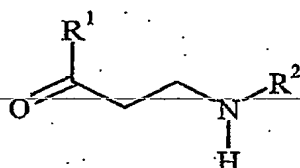
(S,S,S,S)-Me-KetalPhos



(S) and (R)-MeO-BiPhep

8. The process of any of claims 1 to 7, wherein the compounds of formulae Ia and Ib are obtained from their corresponding salts with a carboxylic acid by hydrolysis in the presence of an alkali- or earth alkali hydroxide.

9. Salts of a carboxylic acid with an aminoketone of the formula

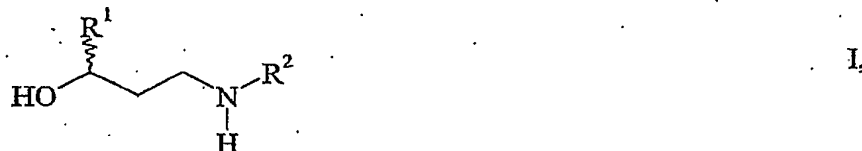


II,

wherein R^1 is selected from the group consisting of 2-thienyl, 2-furanyl and phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups, and wherein R^2 is C_{1-4} -alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups.

10. Salts of formula II, wherein the acid is selected from the group consisting of C_{1-18} -alkanoic acids, (-)-2,3:4,6-di-*O*-isopropylidene-2-keto-L-gulonic acid, (+)-2,3:4,6-di-*O*-isopropylidene-2-keto-D-gulonic acid, 2-keto-L-gulonic acid, 2-keto-D-gulonic acid, L-aspartic acid, D-aspartic acid, DL-aspartic acid, benzoic acid, 3-methyl-benzoic acid, salicylic acid and 1-, 2- or 3-naphthalene carboxylic acid.

11. Salts of a carboxylic acid with an aminoalkohol of the formula

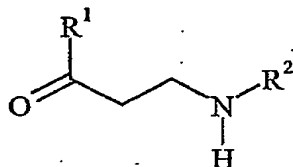


wherein R^1 is selected from the group consisting of 2-thienyl, 2-furanyl and phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups, and wherein R^2 is C_{1-4} -alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups, with the exception of salts, wherein the acid is (-)-2,3:4,6-di-*O*-isopropylidene-2-keto-L-gulonic acid or (+)-2,3:4,6-di-*O*-isopropylidene-2-keto-D-gulonic acid.

18

Abstract

Provided is a process for the preparation of enantiomerically pure 1-substituted-3-amino-alcohols, particularly of (S)-(-)- and (R)-(+)-3-*N*-methylamino-1-(2-thienyl)-1-propanol, by asymmetrically hydrogenating salts of a carboxylic acids with an aminoketone of the formula



wherein R^1 is selected from the group consisting of 2-thienyl, 2-furanyl and phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups, and wherein R^2 is C_{1-4} -alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more C_{1-4} -alkyl or C_{1-4} -alkoxy groups, and wherein the corresponding aminoalcohols are obtained by subsequent hydrolysis of their salts. Furthermore provided are salts of a carboxylic acid with said aminoketones and the aminoalcohols obtained by asymmetrically hydrogenating said aminoketones, respectively.

Basel, April 28, 2004

Dr. Bernd Gallasch